

Amendment and Response to Restriction Requirement
U.S. Application No. 10/509,622

Q8:855

AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application:

LISTING OF CLAIMS:

1. **canceled**
2. **(original):** A polypeptide consisting of the amino acid sequence represented by SEQ ID NO:2.
3. **(currently amended):** A polynucleotide coding for the polypeptide according to claim 1 or claim 2.
4. **(original):** An expression vector comprising the polynucleotide according to claim 3.
5. **(original):** A cell transformed with the expression vector according to claim 4.
6. **(original):** A method for inspecting rheumatoid arthritis, comprising (1) a step of measuring an expression level in a subject of
 - i) a gene comprising the nucleotide sequence according to claim 3, or
 - ii) a gene comprising a nucleotide sequence of a polynucleotide coding for a polypeptide which comprises an amino acid sequence having 95% or more of homology with the amino acid sequence represented by SEQ ID NO:2 and which is expressed specifically in RA patients, and(2) a step of comparing it with an expression level of the gene in a healthy person.
7. **(original):** A rheumatoid arthritis inspection kit which comprises forward and reverse primers designed to specifically amplify
 - i) a gene comprising the nucleotide sequence according to claim 3, or

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ii) a gene comprising a nucleotide sequence of a polynucleotide coding for a polypeptide which comprises an amino acid sequence having 95% or more of homology with the amino acid sequence represented by SEQ ID NO:2 and which is expressed specifically in RA patients.

8. **(currently amended):** A method for screening a substance capable of inhibiting activity of a polypeptide, comprising (1) a step of allowing a substance to be tested to contact with a cell expressing the polypeptide according to ~~claim 1 or claim 2~~ or a polypeptide which comprises an amino acid sequence having 95% or more of homology with the amino acid sequence represented by SEQ ID NO:2 and which is expressed specifically in RA patients, (2) a step of analyzing whether or not activity of the polypeptide is inhibited, and (3) a step of selecting a substance capable of inhibiting activity of the polypeptide.

9. **(currently amended):** The screening method according to claim 8, wherein the substance which inhibits the activity of the polypeptide according to ~~claim 1 or claim 2~~, or of a polypeptide which comprises an amino acid sequence having 95% or more of homology with the amino acid sequence represented by SEQ ID NO:2 and which is expressed specifically in RA patients is a substance for the treatment of rheumatoid arthritis and/or a substance for the treatment of osteoarthritis.

10. **(original):** A method for producing a pharmaceutical composition for the treatment of RA and/or the treatment of osteoarthritis, comprising a step of carrying out screening with the use of the screening method according to claim 8 or claim 9, and a step of formation using a substance obtained by the screening.